

Scientific & Regulatory Consultants, Inc.

December 10, 2015

Document Processing Desk (DCI/AD)
Attn: Rose Kyprianou, Reevaluation Team Leader, PM 36
US EPA (7510P)
One Potomac Yard (South Building)
2777 South Crystal Drive
Arlington, VA 22202

SUBJECT: GDCI-043901-30859 (EPA Co. No. 71814)

Dear Ms. Kyprianou:

On behalf of SteriMed Medical Waste Solutions, Inc. (SteriMed) and in response to the subject DCI, I wish to inform you that SteriMed intends to comply with the requirements set forth in the Notice to maintain their registration as indicated on the attached Data Call In Notice Response.

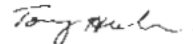
SteriMed is claiming a Generic Data Exemption for their registration pertaining to the subject DCI. SteriMed purchases an EPA registered source of the active ingredient as indicated on the enclosed form. The form has been signed Hilary Newman, Manager of Regulatory & Environmental Affairs at SteriMed. She may be reached by phone at (248) 469-8786.

Submitted with this action:

1. Cover letter
2. Data Call In Response - GDCI-043901-30859

Please contact me at therber@srcconsultants.com or 260-244-6270 if any additional information is needed regarding this response.

Sincerely,



Tony Herber
Agent for SteriMed Medical Waste Solutions, Inc.
cc: Hilary Newman, SteriMed Medical Waste Solutions, Inc.

United States Environmental Protection Agency Washington, D.C. 20460 DATA CALL-IN RESPONSE				OMB Approval 2070-174	
INSTRUCTIONS: Please type or print in ink. Please read carefully the attached instructions and supply the information requested on this form. Use additional sheet(s) if necessary.					
1. Company Name and Address STERIMED MEDICAL WASTE SOLUTIONS, INC. 23065 COMMERCE DR. FARMINGTON HILLS, MI 48335		2. Case # and Name 2315 Glutaraldehyde Chemical # and Name 043901 Glutaraldehyde		3. Date and Type of DCI and Number 28-Aug-2015 GENERIC ID # GDCL-043901-30859	
4. EPA Product Registration	5. I wish to cancel this product registration voluntarily	6. Generic Data		7. Product Specific Data	
		6a. I am claiming a Generic Data Exemption because I obtain the active ingredient from the source EPA registration number listed below.	6b. I agree to satisfy Generic Data requirements as indicated on the attached form entitled "Requirements Status and Registrant's Response."	7a. My product is an MUP and I agree to satisfy the MUP requirements on the attached form entitled "Requirements Status and Registrant's Response."	7b. My product is an EUP and I agree to satisfy the EUP requirements on the attached form entitled "Requirements Status and Registrant's Response."
71814-1				N.A.	N.A.
8. Certification I certify that the statements made on this form and all attachments are true, accurate, and complete. I acknowledge that any knowingly false or misleading statement may be punishable by fine, imprisonment or both under applicable law.				9. Date Signature and Title of Company's Authorized Representative <i>Anthony Nunez</i> Manager of Regulatory & Environmental Affairs	
10. Name of Company SteriMed Medical Waste Solutions, Inc.				11. Phone Number 248.469.8786	